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**(54) Ocular iontophoretic apparatus handle**

Handgriff für eine iontophoretische Augenvorrichtung

Manche destiné à un appareil oculaire à ionophorèse

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• **Koss, Alexander K.**  
**Salt Lake City, Utah 84124 (US)**

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(74) Representative: **Ostertag, Reinhard**  
**Patentanwälte**  
**Dr. Ulrich Ostertag**  
**Dr. Reinhard Ostertag**  
**Eibenweg 10**  
**70597 Stuttgart (DE)**

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(73) Proprietor: **IOMED, INC.**  
**Salt Lake City, Utah 84104 (US)**

(56) References cited:  
**EP-A- 0 927 560 WO-A-99/40967**  
**GB-A- 2 177 928 US-A- 5 169 384**

(72) Inventors:  
• **Beck, Jon E.**  
**Salt Lake City, Utah 84109 (US)**

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## Description

### BACKGROUND OF THE INVENTION

#### 1. Field of the Invention

[0001] The present invention is directed to ocular iontophoretic apparatuses, and more particularly, to a handle for an ocular iontophoretic apparatus to facilitate the grasping, positioning and placement thereof.

#### 2. Background Art

[0002] The use of ocular iontophoretic devices has been known in the art. Such devices as, for example, described in WO-A-99/40967, have been used in an attempt to administer a drug through an electromotive force which drives ionic chemicals through the eye tissue so that they can be absorbed by adjacent tissues and blood vessels.

[0003] Among other problems, difficulties can be incurred with the placement of these devices onto the surface of the eye of a patient. Specifically, inasmuch as certain of these devices are intended for use on only a portion of the eye, they are rather small in size. Accordingly, it is difficult for a doctor to carefully position the device in the proper orientation. Moreover, once placed on the patient's eye, it is often difficult to reposition or to adjust the positioning of the device.

[0004] Accordingly, it is an object of the invention to provide for a handle member which facilitates the grasping, positioning and placement of ocular iontophoretic apparatuses.

[0005] It is likewise an object of the invention to facilitate the repositioning of an ocular iontophoretic apparatus after placement onto the surface of the eye.

[0006] It is a further object of the invention to provide for a handle member which can be pinched so as to flex the iontophoretic apparatus during placement onto an eye and during removal from the eye of the patient.

[0007] These and other objects of the invention will become apparent in light of the specification and claims appended hereto.

### SUMMARY OF THE INVENTION

[0008] The application comprises an iontophoretic apparatus which includes a housing member, a current distribution member, a medicament containment member and a handle member. The current distribution member is associated with the housing member. The medicament containment member is associated with the current distribution member. The handle member is associated with the housing member, and the handle member serves to facilitate the positioning and/or placement of the iontophoretic apparatus.

[0009] The handle member comprises a first handle region and a second handle region extending outwardly

from the handle member. Each of the first and second handle regions are preferably co-molded with the housing member and distally spaced apart a predetermined distance. In one such embodiment, the first and second handle regions extend away from each other. In another embodiment, the handle member may be releasably associated with the housing member.

[0010] In another preferred embodiment, at least one of the first and second handle regions includes a gripping region. The gripping region facilitates the overall gripping of the handle region by a user during the positioning and/or placement of the apparatus. In another such embodiment, the first and second handle regions include a grasping region which joins the handle regions together to, in turn, render a single unitary handle region.

[0011] In another preferred embodiment, the first and second handle regions include means for flexing the housing member. In one such embodiment, the first and second handle regions further include means for limiting the flexing of the housing member. In one embodiment, the flexing limiting means may comprise the positioning of a portion of the first and second handle regions in a spaced apart orientation. The spaced apart orientation substantially corresponds to the desired maximum flex of the housing member.

[0012] In another such embodiment, the flexing limiting means further includes means for aligning the first and second handle regions. The aligning means precludes inadvertent misalignment of the first and second handle regions during flexing.

[0013] In another preferred embodiment, the apparatus further includes means for maintaining the registered placement of the apparatus in the desired orientation. In one such embodiment, the registered placement maintaining means may comprise a receiving region associated with one or both of the handle member and the housing member. In one such embodiment, the receiving region comprises at least one notch. Preferably, the apparatus may further include means for biasing at least a portion of the receiving region against the soft tissue of a user.

### BRIEF DESCRIPTION OF THE DRAWINGS

[0014]

Fig. 1 of the drawings is a perspective view of the first embodiment of the apparatus of the present invention;

Fig. 2 of the drawings is a perspective view of the first embodiment of the apparatus of the present invention;

Fig. 3 of the drawings is a perspective view of the first embodiment of the apparatus of the present invention;

Fig. 4 of the drawings is a perspective view of the first embodiment of the apparatus of the present in-

vention;

Fig. 5 of the drawings is a perspective view of the first embodiment of the apparatus of the present invention;

Fig. 6 of the drawings is a perspective view of the first embodiment of the apparatus of the present invention;

Fig. 7 of the drawings is a perspective view of a second embodiment of the apparatus of the present invention;

Fig. 8 of the drawings is a perspective view of a third embodiment of the apparatus of the present invention;

Fig. 9 of the drawings is a perspective view of a fourth embodiment of the apparatus of the present invention; and

Fig. 10 of the drawings is a perspective view of the fourth embodiment of the apparatus of the present invention.

#### BEST MODE FOR PRACTICING THE INVENTION

[0015] While this invention is susceptible of embodiment in many different forms, there is shown in the drawings and will be described in detail, several specific embodiments with the understanding that the present disclosure is to be considered as an exemplification of the principles of the invention and is not intended to limit the invention to the embodiments illustrated.

[0016] Iontophoretic apparatus 10 is shown in Figs. 1 and 2 as comprising housing member 12, current distribution member 14, medicament containment member 16 and handle member 18. The housing member includes outer surface 40, first end 42, second end 44, upper region 45 and rim 47. Generally, the housing comprises a plastic material which is molded into a desired configuration for the positioning thereof on the eye of a patient.

[0017] As will be understood, housing member 12 may be of any number of sizes and shapes. Various embodiments of the housing member may include various configurations depending on the medicament to be dispensed, as well as the specific shape of the soft tissue surrounding the eye of the patient, and the particular region of the eye to which it is to be applied. Of course, the handle member is not limited to any particular housing member configurations and may be used with a wide variety of such devices. Additionally, the medicament that is retained in medicament containment member 16 for dispensing is not limited to any particular medicament, and virtually any medicament that can be applied iontophoretically through the eye can be used in association with the iontophoretic apparatus.

[0018] Handle member 18 is shown primarily in Figs. 1 and 5 as comprising first handle region 20, second handle region 22, means 24 for flexing the handle member, means 26 for limiting the flexing of the handle member and means 25 for maintaining registered placement

of the apparatus. Generally, handle member 18 is co-molded with housing member 12, however, it is likewise contemplated that the handle member may comprise a separate component which may be welded, adhered or otherwise joined to housing member 12. In addition, the handle member is generally associated with the upper region 45, which, in turn, facilitates placement of the device along, for example, the lower edge of the patient's eye under the lower eyelid. In addition, such handle positioning minimizes the intrusiveness of the handle member and the discomfort to the patient receiving treatment, while maximizing the versatility thereof.

[0019] First handle region 20 of the handle member is shown in Figs. 3 and 4 as including first end 46, second end 48 and first gripping portion 50. First end 46 is associated with outer surface 40 proximate first end 42 of the housing member. Second end 48 extends outwardly therefrom and in a direction which is generally away from second handle region 22 of the handle member. First gripping portion 50 is positioned proximate second end 48 of first handle region 20. Generally, first gripping portion 50 comprises a region which is sized and shaped so as to promote the gripping thereof by a doctor or other professional during placement of the iontophoretic apparatus in the eye of a patient. As can be seen in Figs. 5 and 6, first gripping portion 50 is substantially planar and spaced apart from the housing member a distance sufficient to insure that the doctor can easily grip the gripping portion without inadvertently striking or touching the patient. In certain embodiments the gripping region comprises a substantially flat pod region 51 (Fig. 7).

[0020] It will be understood that second handle region 22 is substantially similar to first handle region 20, and comprises first end 52, second end 54 and second gripping portion 56. While various configurations are contemplated, generally, the first and second handle regions are substantial mirror images of each other (and are substantially symmetrical) about central axis 101 (Fig. 2) of the housing member.

[0021] In the embodiment shown in Fig. 8, gripping portion 50 of first handle region 20 and gripping portion 56 of second handle region 22 are integrally associated with each other, to, in turn, define a single loop like configuration for the gripping portion. In such an embodiment, the integrated gripping portion extends from second end 48 of first handle region 20 to second end 54 of second handle region 22. The integrated gripping portion is substantially parallel to the housing member and generally follows the surface configuration of the outer surface 40 of housing member 12.

[0022] In the embodiment shown in Figs. 9 and 10, handle member 18 may be removably associated with housing member 12. Specifically, in such an embodiment, handle member 18 includes means 74 for releasably retaining the handle member to the housing member. Releasable retaining means 74 includes hoop 76 which is releasably positionable within grooved recess

78 of the housing member. As will be understood, once the apparatus is positioned as desired in the eye, the handle member can be removed by pulling thereon, to, in turn, release hoop 76 from within recess 78 of the housing member. Additionally, it will be understood that while the removable handle member is shown as including a single gripping region, other handle members, including, but not limited to, those shown in Figs. 1, 7 and 8 are likewise contemplated.

[0023] In the embodiment shown in Figs. 5 and 6, gripping portion 50 of first handle region 20 and gripping portion 56 of second handle region 22 are spaced apart a predetermined distance from each other, to essentially provide a means for flexing the housing. Specifically, and as will be explained in more detail below with respect to the operation, as the doctor or professional pinches the first and second gripping portions 50, 56, about the respective second ends of handle regions 20, 22, toward each other, the force, in turn, flexes the housing member. By flexing the housing member during insertion and positioning on the surface of the eye, an improved fit, and an improved positioning can be achieved. Indeed, as long as the first and second gripping portions are not united to form a single integrated gripping portion, by positioning the first and second handle regions at opposite ends of the housing member, the pinching thereof will facilitate the flexing of the housing member.

[0024] Placement of a gap between the first and second gripping portions defines means 26 for limiting the flexing of the housing member. Specifically, gripping portion 50 and gripping portion 56 extend from the respective second ends of the respective handle regions so as to be substantially parallel to outer surface 40 of housing member 12. The two gripping portions essentially extend toward each other until end 62 of first gripping portion 50 is separated from end 64 of second gripping portion 56 by a gap. Thus, as the user pinches the gripping members, the distance separating the two gripping portions becomes smaller until end 62 of first gripping portion is in abutment with end 64 of second gripping portion 56. At such time, the respective gripping portions can be pinched no further and additional flexing of the housing member is not possible.

[0025] As shown in Figs. 5 and 6, the flexing limiting means further includes means 32 for aligning the respective ends of the gripping portions. In particular, aligning means 32 includes first alignment member 68 which is associated with end 60 of gripping portion 50 and second alignment member 70 which is associated with end 62 of gripping portion 52. As will be understood, as the user flexes the ends, the alignment members insure that end 60 of gripping portion 50 is aligned with end 62 of gripping portion 52, so that the flexing can be limited by the interaction and abutment of the two ends. Indeed, without the alignment members, inadvertent misalignment of the ends would permit the flexing of the respective ends without limitation and beyond that which is desirable.

[0026] Placement registration means 25 is shown in Fig. 1 as including retaining regions 70, 72 and means 73 for biasing the retaining region against the soft tissue of a patient. Retaining region 70 is associated with one or both of first end 46 of first handle region 20 and first end 42 of housing member 12. Similarly, retaining region 72 is associated with one or both of first end 52 of second handle region 22 and second end 44 of housing member 12. The retaining regions comprise structures, such as notches that are configured to cooperate with the corner area of the soft tissue surrounding the eye of the user. Biasing means 73 comprises the natural resilience of the material surrounding retaining region 70 to return to its original configuration upon flexing thereby biasing the retaining regions against the soft tissue of a patient. As will be explained, the corner areas of the eye, the retaining regions and the biasing means cooperate to maintain the registered placement of the apparatus in the desired orientation, and substantially preclude movement of the apparatus once positioned. It is also contemplated that the placement registration means is not limited to cooperation with the corners of the eyes; rather, it can also cooperate with other surrounding tissues that remain substantially static during movement of the eye.

[0027] It is additionally contemplated that the electrical leads which attach the power supply to the electrodes which drive the medicament may be molded into the handle member.

[0028] In operation, the doctor, physician's assistant or other professional first selects the appropriate apparatus from among various apparatuses of different size, shape and medicament. As explained above, the apparatus is not limited to any particular shape and any particular medicament. Once selected and prepared for placement by the doctor or assistant on the patient's eye, the apparatus is grasped by the gripping members and positioned onto the surface of the eye.

[0029] In particular, in the embodiment shown in Fig. 1, the doctor first pinches the respective gripping portions 50, 56 toward each other so as to flex housing member 12. Once flexed as desired, the user positions the housing member on the surface of the patient's eye. As the initial contact with the surface of the eye is attained, the doctor slowly releases the pinching grip on the gripping regions, and housing member 12 returns to its original orientation at which time the entire outer rim 46 is in contact with the surface of the eye. By pinching the gripping regions prior to positioning, improved surface mating between the outer rim and eye and improved comfort to the patient is achieved. Moreover, better control can be maintained over the apparatus, which, in turn, facilitates improved accuracy relative to placement on the surface of the eye.

[0030] In addition, as the doctor releases the handle member, the corner regions of the eye are positioned into and accepted by retaining regions 70 and 72 of registered placement maintaining means 25 so as to

achieve registered placement of the apparatus in the desired orientation.

[0031] Furthermore, biasing means 73 of registered placement maintaining means 25 cooperates with the retaining regions to bias and, in turn, maintain the corner of the eye and the retaining regions in cooperative engagement. The cooperation of the registered placement maintaining means and the biasing means can also compensate for size variation in the dimensions of the soft tissue surrounding the eye. In certain embodiments, biasing means 73 and the flexing means of the handle member may be integrated into a single structure.

[0032] Once fully positioned, the doctor initiates current delivery from the current distribution member. The current forces medicament retained in the medicament containment member through the tissue of the patient's eye. The treatment continues for a predetermined period of time which is determined by the type and quantity of medicament that is to be transmitted to the patient.

[0033] Once the treatment is complete, current ceases to be delivered by the current distribution member. At such time, passage of medicament through the patient's tissue ceases. When the treatment is complete, the apparatus can be removed from the patient. Specifically, the doctor again grasps the gripping regions of handle member 12 and pulls the housing from the surface of the eye.

[0034] In the embodiment shown in Fig. 1, the user pinches gripping portions 50 and 56 until housing member 12 flexes. The flexing of the housing member likewise facilitates the release of the housing member from the surface of the eye in a controlled manner.

[0035] In the embodiment shown in Fig. 7, the user may individually grasp each of the separate gripping members to release the housing member from the surface of the eye. Similarly, in the embodiment of Fig. 8, the doctor can grab the unitized gripping member to release the housing member from the surface of the eye.

[0036] The foregoing description merely explains and illustrates the invention and the invention is not limited thereto except insofar as the appended claims are so limited, as those skilled in the art who have the disclosure before them will be able to make modifications without departing from the scope of the invention as defined in the claims.

## Claims

1. An ocular iontophoretic apparatus comprising:

- a housing member (12);
- a current distribution member (14) associated with the housing member (12);
- a medicament containment member (16) associated with the current distribution member (14); and
- a handle member (18) associated with the

housing member (12), the handle member (18) facilitating the placement, positioning, registration and securement of the iontophoretic apparatus,

### characterized in that

- the handle member (18) comprises a first handle region (20) and a second handle region (22) extending outwardly from the handle member (18), the first and second handle regions (20, 22) distally spaced apart a predetermined distance.

2. The apparatus of claim 1, **characterized in that** at least one of the first and second handle regions (20, 22) includes a gripping region (50, 56), the gripping region (50, 56) facilitating the grasping of the handle region (20, 22) by a user.

3. The apparatus of claim 1, **characterized in that** the first and second handle regions (20, 22) extend away from each other.

4. The apparatus of claim 1, **characterized in that** the first and second handle regions (20, 22) include a grasping region (50, 56) which joins the handle regions (20, 22) into a single unitary handle region.

5. The apparatus of claim 1, **characterized in that** the first and second handle regions (20, 22) include means (24) for flexing the housing member (12).

6. The apparatus of claim 5, **characterized in that** the first and second handle regions (20, 22) include means (26) for limiting the flexing of the housing member (12).

7. The apparatus of claim 6, **characterized in that** flexing limiting means (26) comprises the positioning of at least a portion of each of the first and second handle regions (20, 22) in a spaced apart orientation, substantially corresponding to the desired maximum flex of the housing member (12).

8. The apparatus of claim 7, **characterized in that** the flexing limiting means (26) further includes means (32) for aligning the first and second handle regions (20, 22) to, in turn preclude inadvertent misalignment thereof.

9. The apparatus of claim 1, **characterized in that** the first and second handle regions (20, 22) are molded with the housing member (12).

10. The apparatus of claim 1, **characterized in that** the handle member (18) is releasably associated with the housing member (12).

11. The apparatus of claim 1, further including means (25) for maintaining the registered placement of the apparatus in the desired orientation.
12. The apparatus of claim 11, **characterized in that** the registered placement maintaining means (25) comprises at least one receiving region (70, 72) associated with at least one of the housing member (12) and the handle member (18), the receiving region (70, 72) configured to cooperate with a portion of the soft tissue of a user.
13. The apparatus of claim 12, **characterized in that** the receiving region (70, 72) comprises at least one notch.
14. The apparatus of claim 12, further including means (73) for biasing at least a portion of the receiving region (70, 72) relative to a portion of the soft tissue of a user, to, in turn, further maintain registered placement thereof within the eye of a user.

#### Patentansprüche

##### 1. Augeniontophoreseapparatur, umfassend:

- einen Gehäuseteil (12);
- ein Stromverteilungselement (14), das mit dem Gehäuseteil (12) verbunden ist;
- ein Medikamentenbehälterelement (16), das dem Stromverteilungselement (14) zugeordnet ist; und
- ein Griffelement (18), das mit dem Gehäuseteil (12) verbunden ist, wobei das Griffelement (18) die Anordnung, Positionierung, die Ausrichtung und die Befestigung der Iontophoreseapparatur erleichtert,  
**dadurch gekennzeichnet, dass**
- das Griffelement (18) einen ersten Griffbereich (20) und einen zweiten Griffbereich (22) umfasst, die sich nach außen von dem Griffelement (18) erstrecken, wobei der erste und der zweite Griffbereich (20, 22) distal mit einer vorbestimmten Entfernung beabstandet sind.

2. Apparatur nach Anspruch 1, **dadurch gekennzeichnet, dass** der erste und/oder zweite Griffbereich (20, 22) eine Greifzone (50, 56) umfassen, wobei die Greifzone (50, 56) das Ergreifen des Griffbereichs (20, 22) durch einen Benutzer erleichtert.
3. Apparatur nach Anspruch 1, **dadurch gekennzeichnet, dass** der erste und der zweite Griffbereich (20, 22) sich voneinander weg erstrecken.
4. Apparatur nach Anspruch 1, **dadurch gekennzeichnet, dass** der erste und der zweite Griffbe-

reich (20, 22) eine Greifzone (50, 56) umfassen, der die Griffbereiche (20, 22) zu einem einzigen ganzheitlichen Griffbereich verbindet.

5. Apparatur nach Anspruch 1, **dadurch gekennzeichnet, dass** der erste und der zweite Griffbereich (20, 22) Mittel (24) zum Biegen des Gehäuseteils (12) umfassen.
6. Apparatur nach Anspruch 5, **dadurch gekennzeichnet, dass** der erste und der zweite Griffbereich (20, 22) Mittel (26) zur Begrenzung des Biegens des Gehäuseteils (12) umfassen.
7. Apparatur nach Anspruch 6, **dadurch gekennzeichnet, dass** die das Biegen begrenzenden Mittel (26) die Anordnung von mindestens einem Teil sowohl des ersten als auch des zweiten Griffbereichs (20, 22) in einer beabstandeten Orientierung umfassen, welche im Wesentlichen der gewünschten maximalen Biegung des Gehäuseteils (12) entspricht.
8. Apparatur nach Anspruch 7, **dadurch gekennzeichnet, dass** die die Biegung begrenzenden Mittel (26) weiter Mittel (32) zur Ausrichtung des ersten und des zweiten Griffbereichs (20, 22) umfassen, um ihrerseits eine versehentliche Fehlausrichtung derselben auszuschließen.
9. Apparatur nach Anspruch 1, **dadurch gekennzeichnet, dass** der erste und der zweite Griffbereich (20, 22) zusammen mit dem Gehäuseteil (12) geformt sind.
10. Apparatur nach Anspruch 1, **dadurch gekennzeichnet, dass** das Griffelement (18) lösbar mit dem Gehäuseteil (12) verbunden ist.
11. Apparatur nach Anspruch 1, weiter umfassend Mittel (25) zur Aufrechterhaltung der passgenauen Anordnung der Apparatur in der gewünschten Orientierung.
12. Apparatur nach Anspruch 11, **dadurch gekennzeichnet, dass** die die passgenaue Anordnung aufrechterhaltenden Mittel (25) mindestens einen Aufnahmebereich (70, 72) umfassen, der dem Gehäuseteil (12) und/oder dem Griffelement (18) zugeordnet ist, wobei der Aufnahmebereich (70, 72) so konfiguriert ist, dass er mit einem Teil des weichen Gewebes eines Benutzers zusammenarbeitet.
13. Apparatur nach Anspruch 12, **dadurch gekennzeichnet, dass** der Aufnahmebereich (70, 72) mindestens eine Kerbe umfasst.
14. Apparatur nach Anspruch 12, weiter umfassend

Mittel (73) zur Vorspannung mindestens eines Teils des Aufnahmebereichs (70, 72) relativ zu einem Teil des weichen Gewebes eines Benutzers, um wiederum weiter eine passgenaue Anordnung derselben innerhalb des Auges eines Benutzers aufrechtzuerhalten.

## Revendications

### 1. Appareil d'ionophorèse oculaire, comprenant :

- un élément formant enveloppe (12) ;
- un élément distributeur de courant (14) associé à l'élément formant enveloppe (12) ;
- un élément formant réservoir à médicament (16), associé à l'élément distributeur de courant (14) ; et
- un élément formant poignée (18) associé à l'élément formant enveloppe (12), l'élément formant poignée (18) facilitant la mise en place, le positionnement, la mise en coïncidence et l'assujettissement de l'appareil d'ionophorèse, **caractérisé en ce que**
- l'élément formant poignée (18) comprend une première région de poignée (20) et une deuxième région de poignée (22) s'étendant vers l'extérieur à partir de l'élément formant poignée (18), la première et la deuxième régions de poignée (20, 22) étant distalement espacées d'une distance prédéterminée.

2. Appareil selon la revendication 1, **caractérisé en ce qu'**au moins l'une de la première et la deuxième régions de poignée (20, 22) comprend une région de préhension (50, 56), la région de préhension (50, 56) facilitant la prise en main, par un utilisateur, de la région formant poignée (20, 22).

3. Appareil selon la revendication 1, **caractérisé en ce que** la première et la deuxième régions de poignée (20, 22) sont à distance l'une de l'autre.

4. Appareil selon la revendication 1, **caractérisé en ce que** la première et la deuxième régions de poignée (20, 22) comprennent une région de prise en main (50, 56), qui réunit en une région de poignée monobloc unique les régions de poignée (20, 22).

5. Appareil selon la revendication 1, **caractérisé en ce que** la première et la deuxième régions de poignée (20, 22) comprennent un moyen (24) pour fléchir l'élément formant enveloppe (12).

6. Appareil selon la revendication 5, **caractérisé en ce que** la première et la deuxième régions de poignée (20, 22) comprennent un moyen (26) pour limiter la flexion de l'élément formant enveloppe (12).

7. Appareil selon la revendication 6, **caractérisé en ce que** le moyen limiteur de flexion (26) comprend le positionnement d'au moins une partie de chacune de la première et la deuxième régions de poignée (20, 22) selon une orientation qui les écarte l'une de l'autre, d'une manière correspondant pour l'essentiel à la flexion maximale souhaitée de l'élément formant enveloppe (12).

8. Appareil selon la revendication 7, **caractérisé en ce que** le moyen limiteur de flexion (26) comprend en outre un moyen (32) pour aligner la première et la deuxième régions de poignée (20, 22), pour de ce fait en empêcher un mésalignement par inadvertance.

9. Appareil selon la revendication 1, **caractérisé en ce que** la première et la deuxième régions de poignée (20, 22) sont moulées en même temps que l'élément formant enveloppe (12).

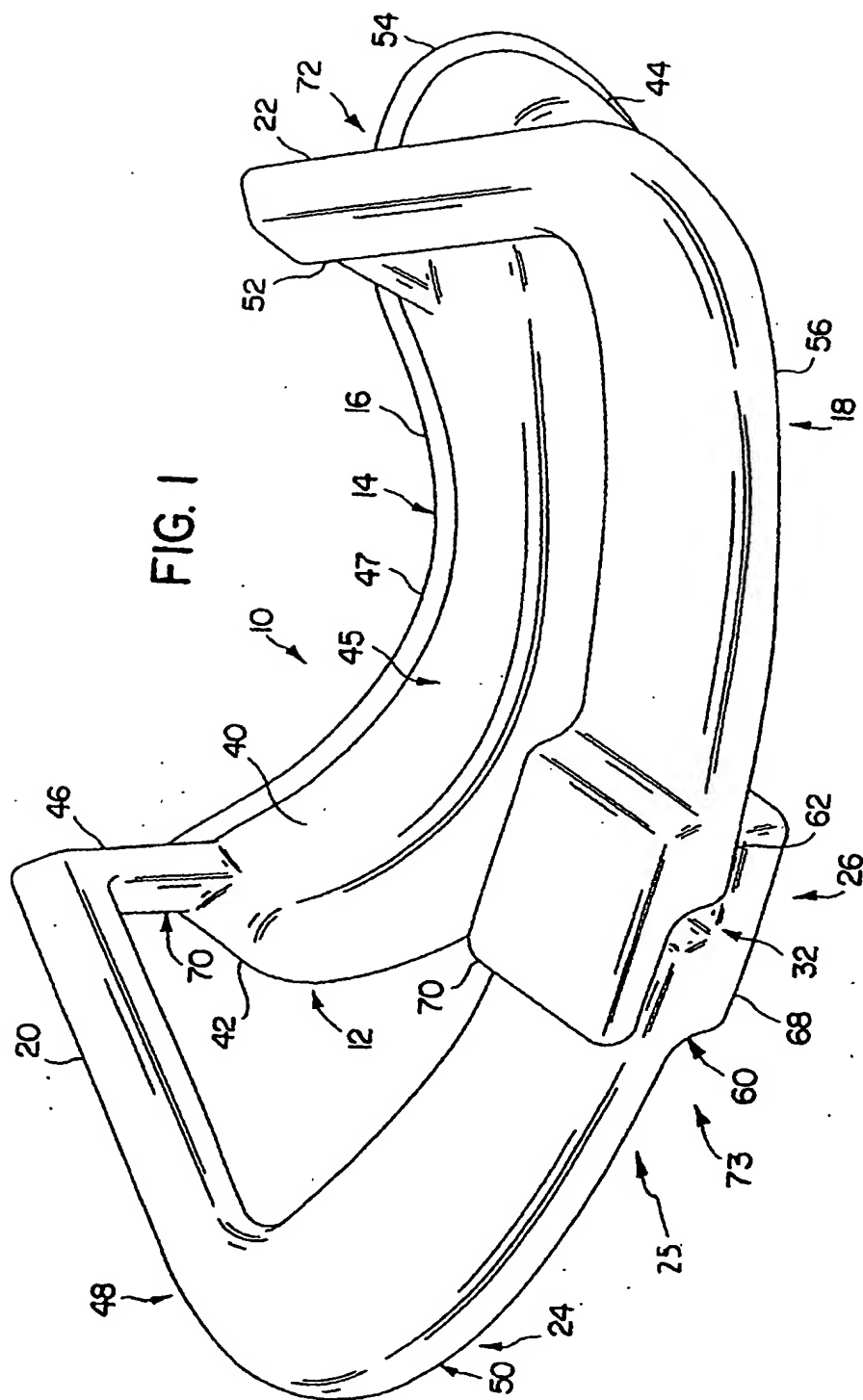
10. Appareil selon la revendication 1, **caractérisé en ce que** l'élément formant poignée (18) est associé d'une manière amovible à l'élément formant enveloppe (12).

11. Appareil selon la revendication 1, qui comprend en outre un moyen (25) pour maintenir en place et en coïncidence l'appareil selon l'orientation voulue.

12. Appareil selon la revendication 11, **caractérisé en ce que** le moyen (25) assurant le maintien de la mise en place en coïncidence comprend au moins une région réceptrice (70, 72) associée à au moins l'un de l'élément formant enveloppe (12) et de l'élément formant poignée (18), la région réceptrice (70, 72) étant configurée de façon à coopérer avec une partie du tissu mou d'un utilisateur.

13. Appareil selon la revendication 12, **caractérisé en ce que** la région réceptrice (70, 72) comprend au moins une encoche.

14. Appareil selon la revendication 12, qui comprend en outre un moyen (73) pour décaler au moins une partie de la région réceptrice (70, 72) par rapport à une partie du tissu mou d'un utilisateur, pour de ce fait maintenir encore plus sa mise en place en coïncidence à l'intérieur de l'oeil d'un utilisateur.





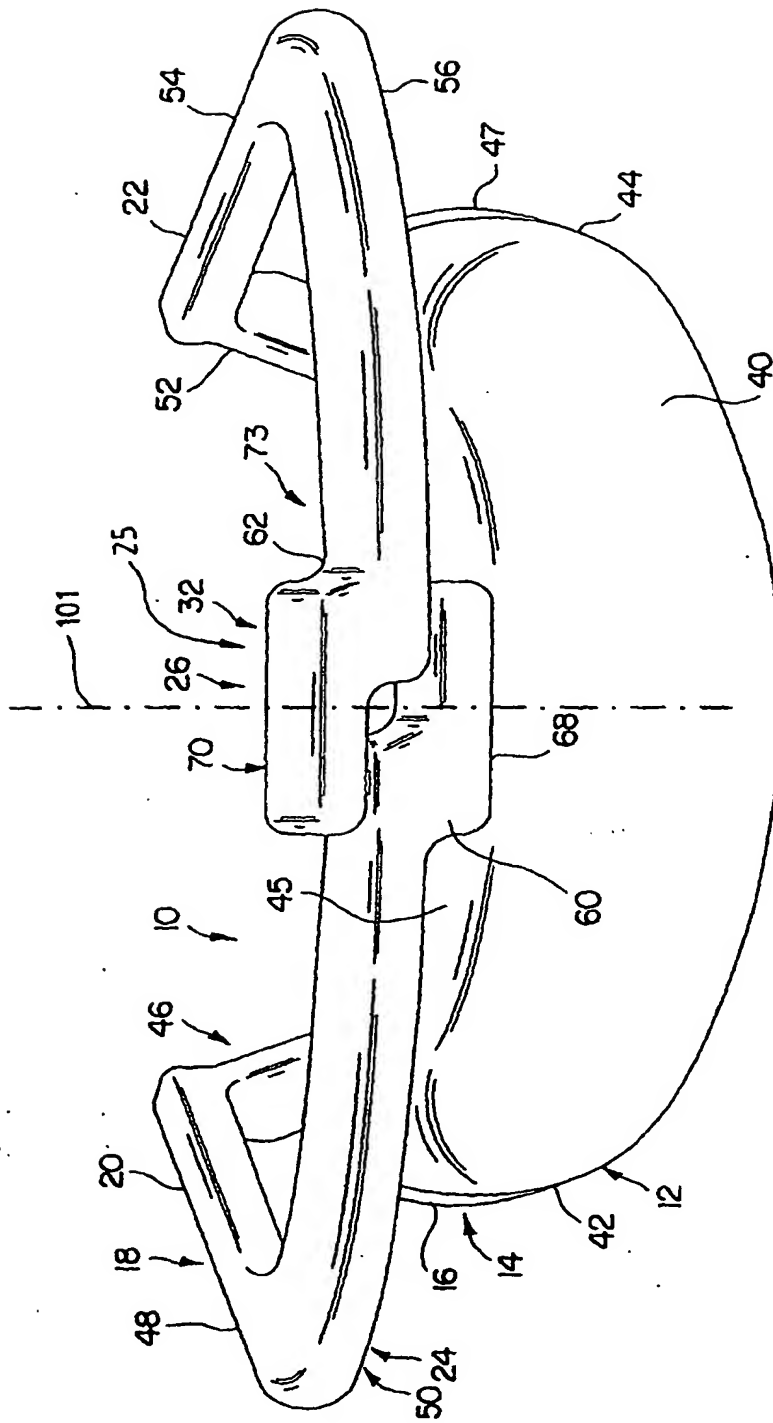
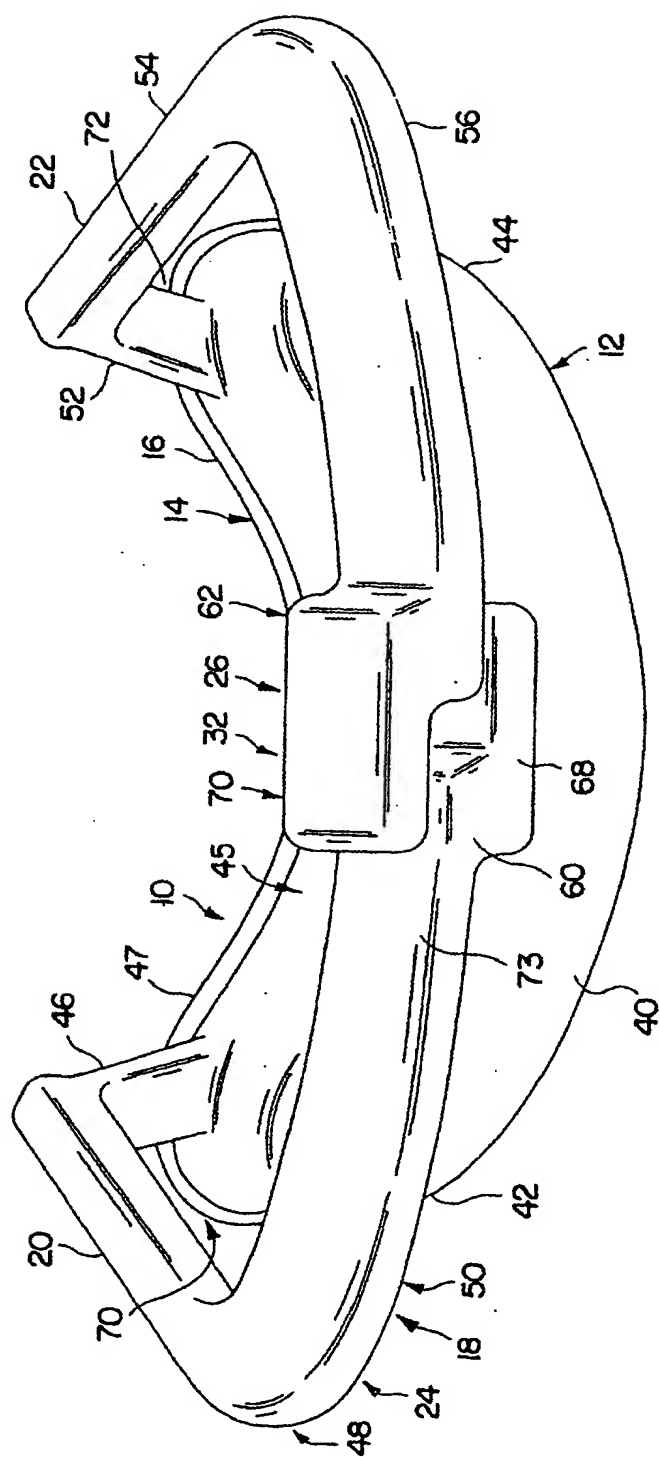


FIG. 2



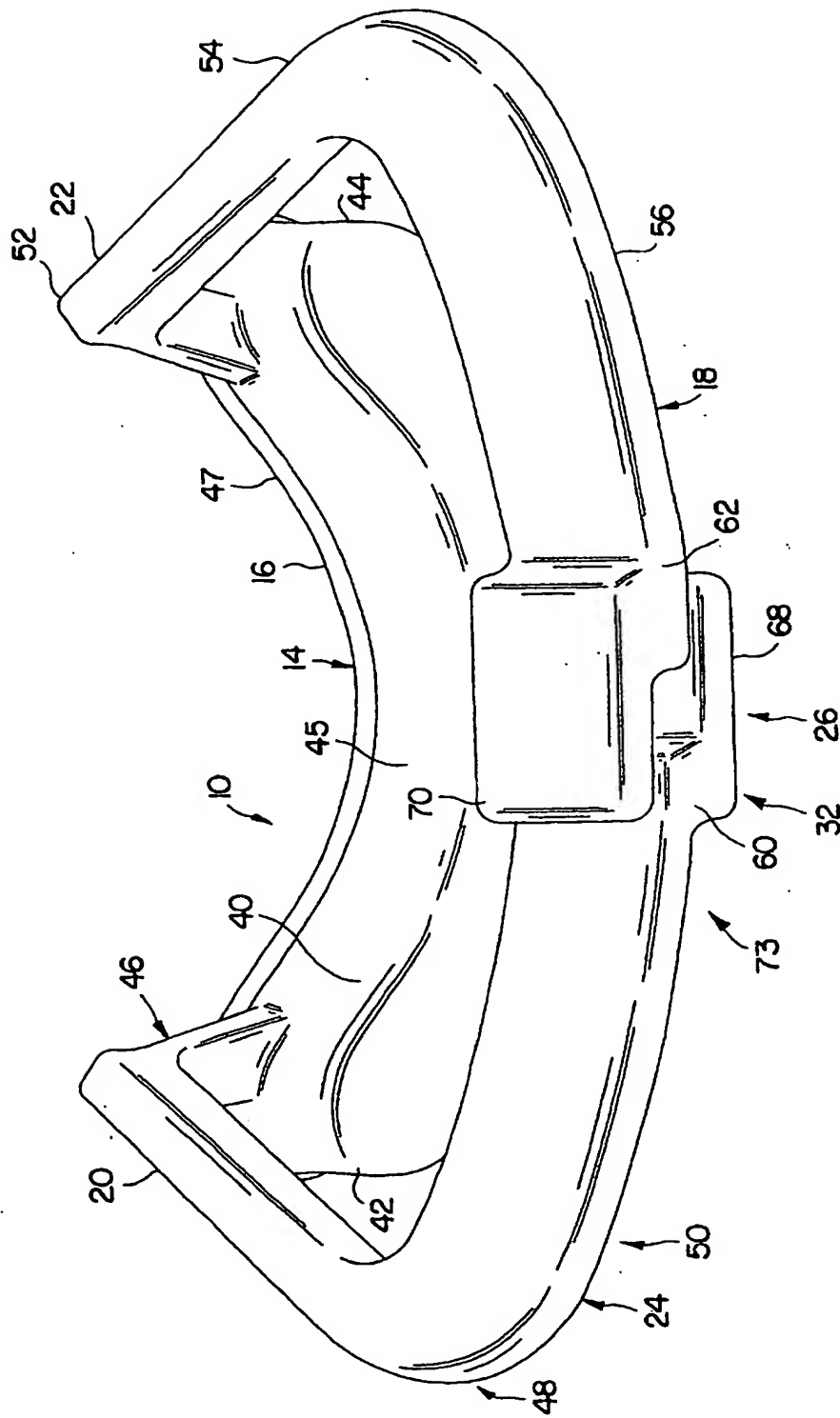
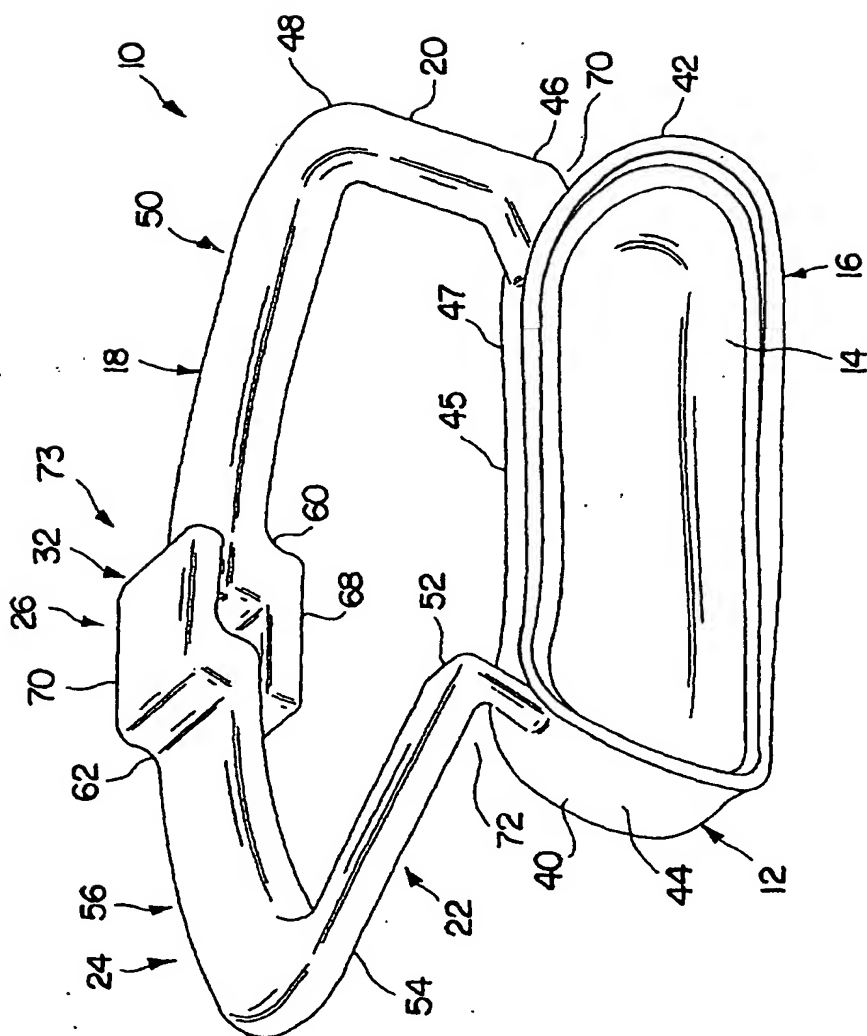
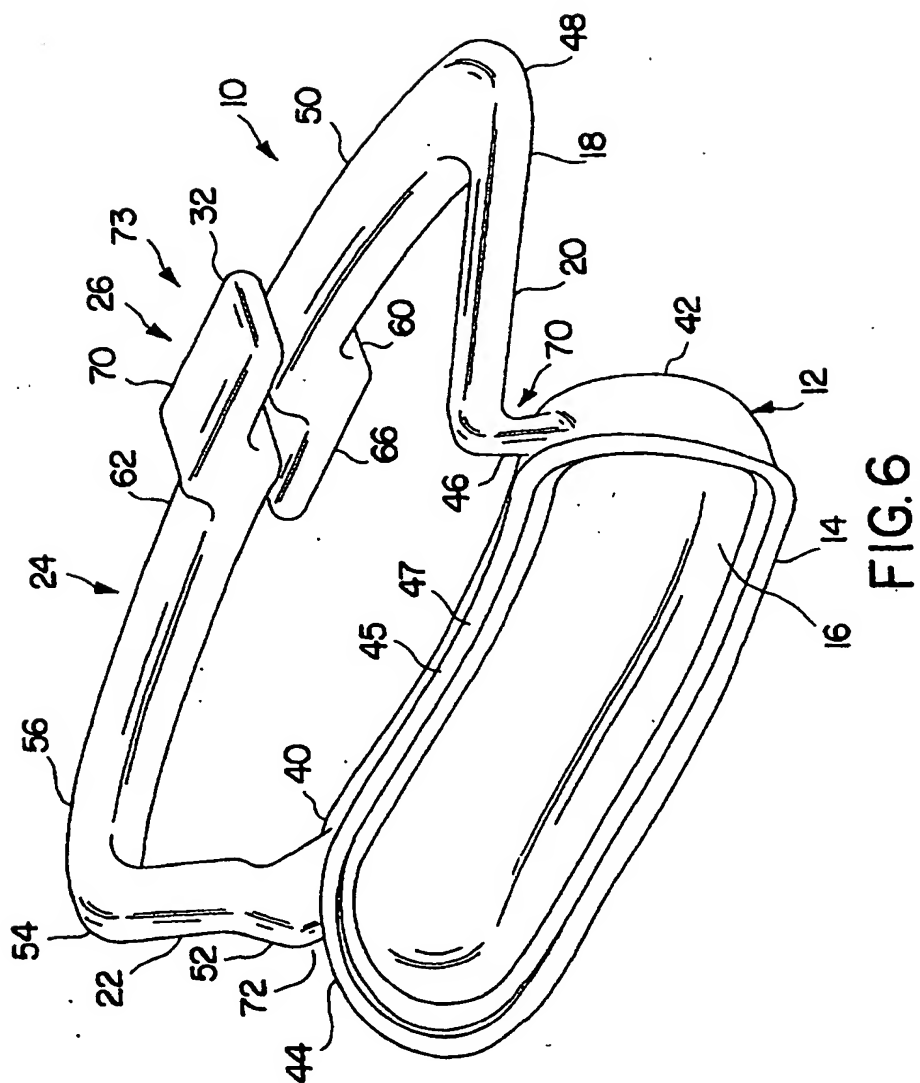
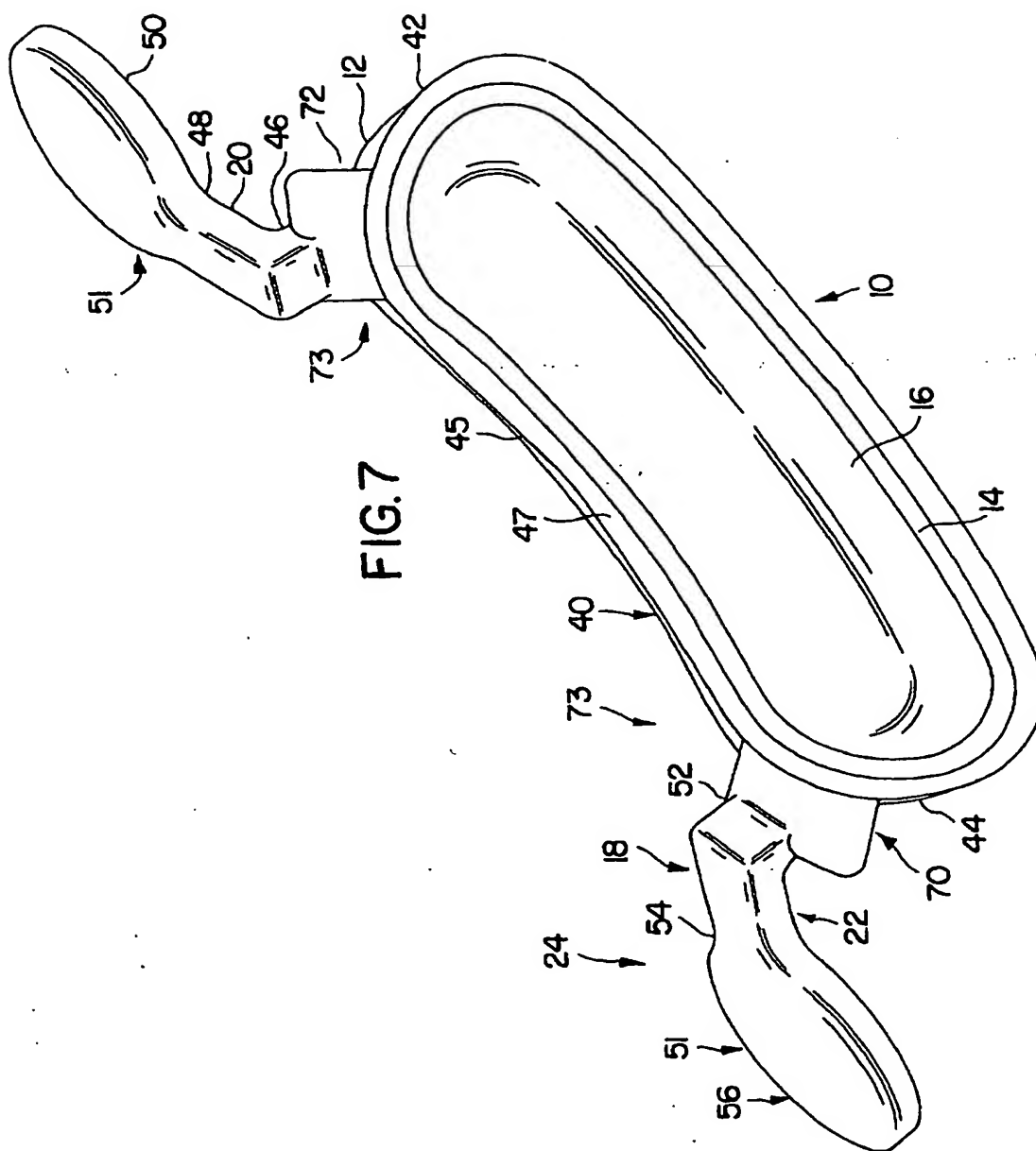


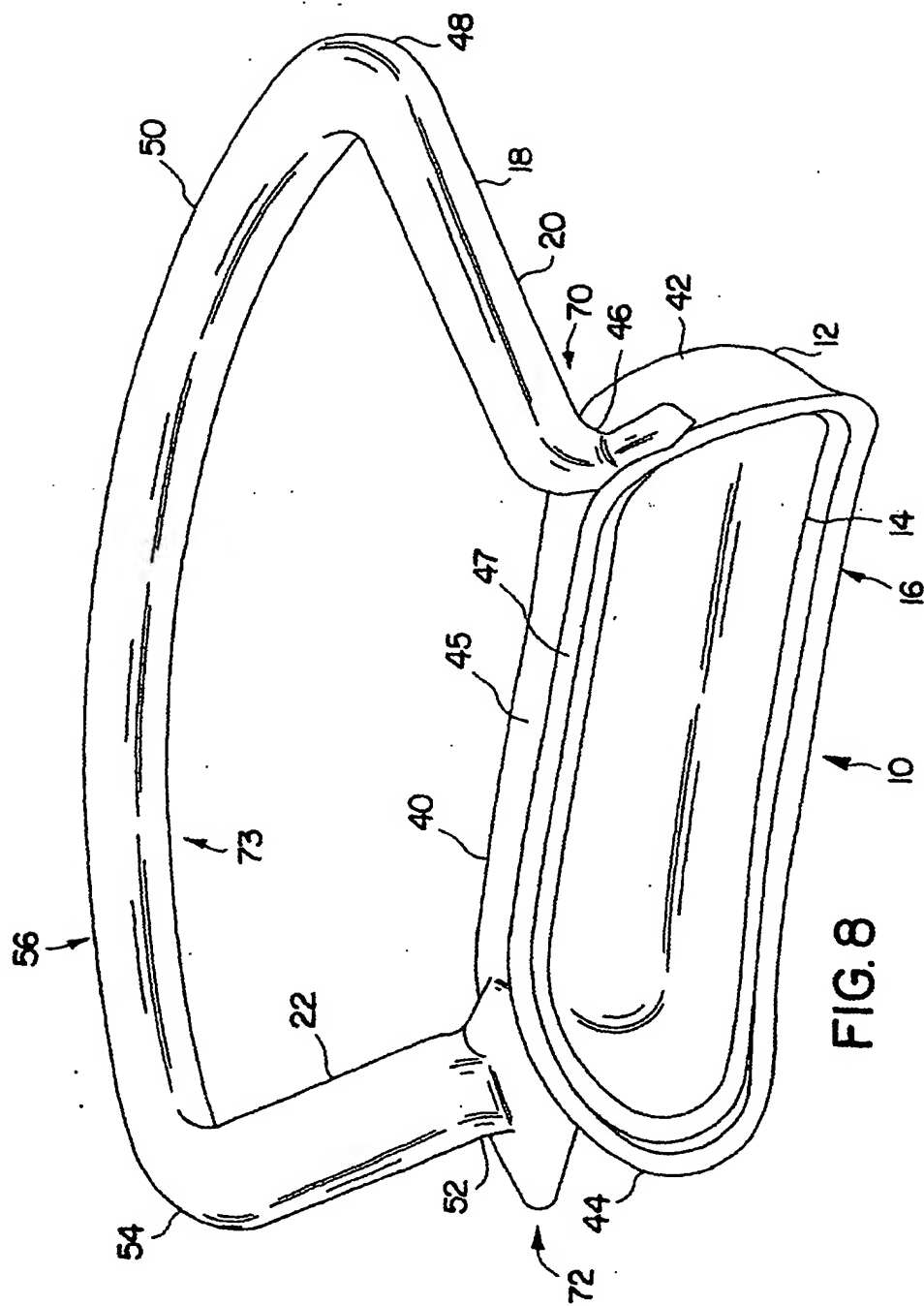
FIG. 4



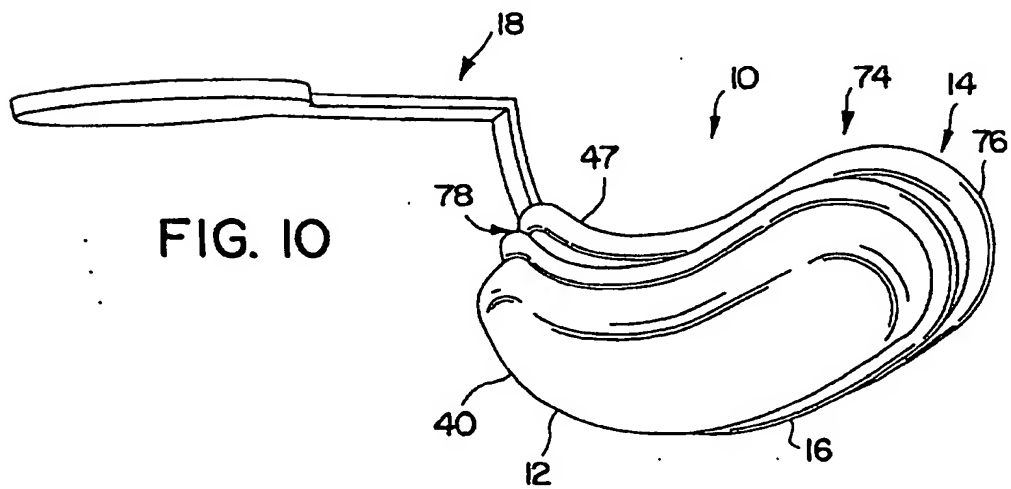
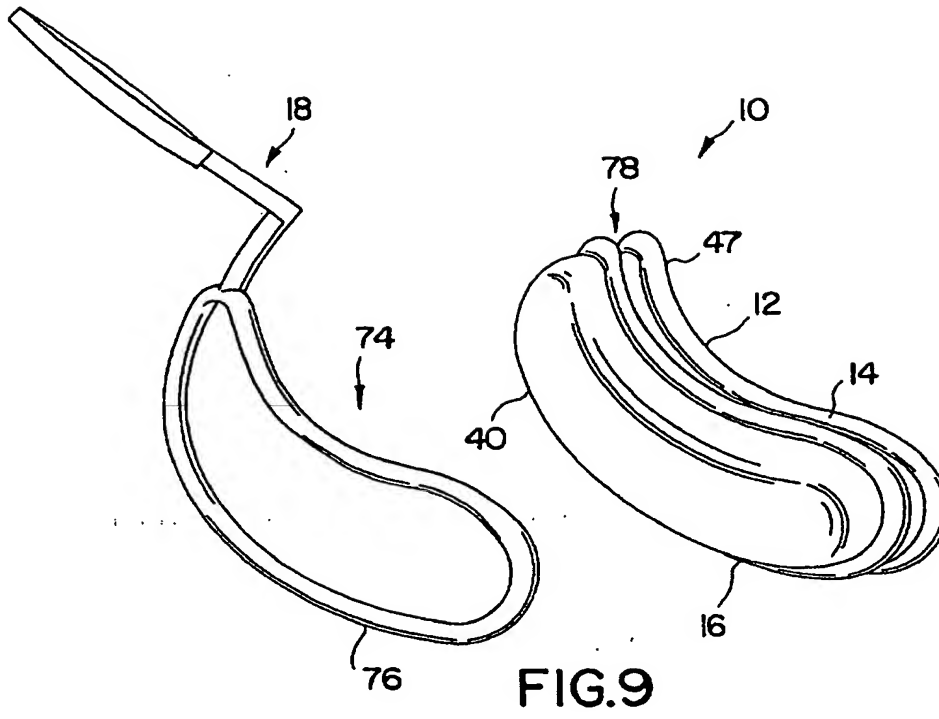
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8/6/7





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